

(19)



Europäisches Patentamt
European Patent Office
Office européen des brevets



(11)

EP 0 776 672 A1

(12)

EUROPEAN PATENT APPLICATION

(43) Date of publication:
04.06.1997 Bulletin 1997/23

(51) Int. Cl.⁶: **A61M 16/00**

(21) Application number: **96116014.0**

(22) Date of filing: **07.10.1996**

(84) Designated Contracting States:
CH DE ES FR GB IT LI NL

(71) Applicant: **Siemens-Elema AB**
171 95 Solna 1 (SE)

(30) Priority: **01.12.1995 SE 9504312**

(72) Inventor: **Nord, Magnus**
161 51 Bromma (SE)

(54) Method for controlling a breathing apparatus and a breathing apparatus

(57) A method for controlling a breathing apparatus is described. In the method, momentary compliance (38, 44, 46) is calculated during an inspiration (40A, 40B). The calculated momentary compliance (38, 44, 46) is then compared to a threshold value (42). If momentary compliance (38) is less than the threshold value (42) during a first interval (40A), positive end expiratory pressure (PEEP) is reduced for subsequent breathing cycles so momentary compliance is greater than the threshold value (42) during the first interval (40A). If momentary compliance (44) is less than the threshold value (42) during a second interval (40B), the ratio between inspiration time and expiration time, as well as the breathing rate, is changed so momentary compliance is greater than the threshold value (42) during the second interval (40B), at the same time as a pre-defined minute volume is generated.

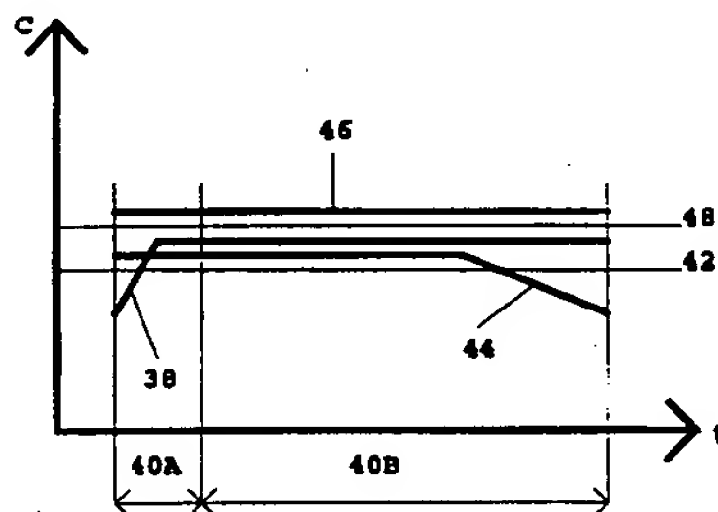


FIG. 3

EP 0 776 672 A1

Description

The present invention relates to a method for controlling a breathing apparatus according to the preamble to claim 1.

Breathing apparatuses supplying a breathing gas to a respiratory system (in humans or animals) and carrying expired breathing gas out of the respiratory system must be controlled in some way in order to avoid risks of damage to the respiratory system. In particular, the supply operation must be appropriately controlled. Preventing an excessive rise in pressure is essential, since excessive pressure could cause barotrauma. In the corresponding manner, supplying large volumes of gas to the respiratory system could cause volutrauma.

This is particularly the case in the ventilation of patients with diseased or damaged lungs. Ventilators connected to the patient's lungs are generally equipped with, or connected to, flow and pressure meters. Pressure and volume in the lungs can thus be monitored with the aid of pressure and flow measurements.

One problem in monitoring a patient with respect to pressure and volume is that damage-causing levels of pressure and volume can vary from patient to patient. In some patients, damaging pressure builds up in certain parts of the lung, whereas pressure remains on a harmless level in other parts of the lung.

At the same time, pressure must not be allowed to drop too much in certain patients, since their lungs might then collapse, making it necessary to supply an extra large amount of breathing gas to re-open the lungs. Lung collapse can also be partial, i.e. only parts of the lungs collapse. Positive end expiratory pressure (PEEP), a greater than atmospheric pressure produced at the end of expiration, is used for keeping the lung open until the following inspiration commences.

In addition, the patient must also be supplied with a sufficient amount of breathing gas. Breathing gas supplied can be designated in terms of the minute volume supplied.

One object of the invention is to achieve a method for controlling breathing apparatuses which solves the afore-mentioned problem.

Another object of the invention is to achieve a breathing apparatus which permits safe and reliable delivery of breathing gas to a respiratory system.

The first object is achieved in accordance with the invention in that the method for controlling the breathing apparatus is devised as set forth in the characterizing part of claim 1.

Compliance designates the elasticity of the respiratory system. It is determined as the ratio between volume and pressure in the respiratory system. A healthy lung has high compliance and can therefore accommodate relatively large changes in volume without major changes in pressure. Pressure rises rapidly, when there is a small increase in volume, only when the lung's physical volume limitations are approached. In other words, compliance drops rapidly when the healthy lung

nears its maximum volume.

The situation is rather different for a damaged or diseased lung. At the start of inspiration with a collapsed lung, pressure rises rapidly with small changes in volume. So compliance is initially very poor. When the lung then opens, compliance is more like the compliance of the healthy lung but is usually much poorer. If the lung is also inherently stiff (atelectatic), the upper limit for volume capacity is reached more quickly, i.e. the stiffness of the lung, rather than the thorax, governs when the lung is full, and compliance drops rapidly. In other words, the damaged or diseased lung has a much smaller effective ventilation range than a healthy lung.

Momentary compliance (the ratio between the derivative of volume and the derivative of pressure in the respiratory system) is therefore an excellent control parameter. When an appropriate threshold value is determined for each patient, parameters such as the level of pressure, PEEP, inspiration time, expiration time, respiratory rate etc. can be automatically controlled by the breathing apparatus.

Advantageous embodiments of the method will be apparent from the dependent claims.

A breathing apparatus is achieved in accordance with the invention with an apparatus devised according to claim 7. Advantageous embodiments are set forth in the dependent claims to claim 7.

One embodiment of the invention will be described below, referring to the figures in which

FIG. 1 shows an embodiment of the breathing apparatus according to the invention;

FIG. 2 shows a respiratory curve and

FIG. 3 shows several compliance curves during inspiration.

In FIG. 1, a breathing apparatus 2 is connected to a patient 4 to supply same with breathing gas and remove expired breathing gas.

The breathing apparatus 2 can accommodate one or a plurality of gases, via three gas connectors 6A, 6B, 6C, which are then mixed into a breathing gas in a mixing chamber 8. Regulation of breathing gas supplied to the patient 4 is performed via an inspiratory valve 10 which is regulated by a control device 12. Alternately, the respective gas can be regulated at the gas connectors 6A, 6B, 6C, i.e. before the breathing gas is mixed in the mixing chamber 8.

Breathing gas is carried from the breathing apparatus 2 to the patient 4 in an inspiratory line 14 and a connector line 16. The connector line 16 can comprise a breathing mask or a Y-piece with a tracheal tube or some other known connector means. Expired breathing gas is carried from the patient 4 back to the breathing apparatus 2 through the connector line 16 and an expiratory line 18. An expiratory valve 20 is arranged in the expiratory line 18. The expiratory valve 20 is regulated by the control device 12. With it a positive end expiratory pressure (PEEP) can be maintained at the end of expi-

ration to e.g. prevent collapse of the patient's 4 lung. Expired breathing gas can be discharged into ambient air through an evacuation unit 22 or collected from the evacuation unit 22 for analysis, filtering or similar.

A pressure meter 24 and a flow meter 26 are arranged in the breathing apparatus 2 to measure the pressure and flow of the breathing gas. The measurement signals can be used for regulating the inspiratory valve 10 so the correct pressure and/or flow are/is supplied to the patient 4. With the aid of measurement signals, pressure and flow in or near the patient's 4 lungs and airways can be calculated. Volume can be determined from flow. Momentary compliance can be determined when pressure and flow are known, as is described in greater detail in comments on FIG. 2. It should be noted that pressure and flow meters can also be arranged near the patient 4, as illustrated with the meters 28, 30 in FIG. 1. More accurate values for the actual conditions in or near the patient 4 are accordingly obtained. The pressure meter can also be situated in the patient's 4 respiratory system, e.g. near carina.

The pressure-volume diagram in FIG. 2 shows an inspiratory and expiratory curve for a diseased or damaged lung. The diagram shows an inspiratory curve 32 and an expiratory curve 34. "Pressure" refers to absolute pressure in the lung, and "volume" refers to the supplied (inspired) volume. As the inspiratory curve 32 shows, pressure initially (area 36A) rises more rapidly than volume. This may be because the lung has collapsed in whole or part and a large positive pressure is needed to open the lung to permit the influx of breathing gas. So compliance in area 36A is accordingly poor.

When the lung has opened (area 36B), breathing gas can flow in more easily, so pressure does not increase as rapidly. This area displays the lung's maximum compliance. The lung ultimately expands as much as it can and is accordingly full. Breathing gas is unable to flow in as easily (area 36C), and every increase in volume causes a sharp rise in pressure. So compliance is again poor in the area 36C.

During expiration (curve 34), pressure and volume drop back to their initial values. Since expiration is passive, and the expiratory valve 20 in the breathing apparatus 2 regulates flow and pressure in expiration, the expiratory curve 34 is therefore of less interest than the inspiratory curve 32.

If momentary compliance is determined, the breathing apparatus can be controlled so ventilation only takes place in area 36B in which the lung displays maximum compliance. Here, momentary compliance can be determined in a plurality of ways. Volume and pressure can be established and the ratio of the respective derivatives can be calculated. Alternately, the momentary flow value can be divided by the derivative of momentary pressure. Derivatives can be determined in the known manner.

FIG. 3 shows three of the situations, which can occur during an inspiration, in order to illustrate the invention. The diagram shows compliance on one axis

and time on the other. A first compliance curve 38 shows that compliance in a first interval 40A rises sharply and passes a first threshold value 42. Compliance then remains constant for the rest of inspiration, a second interval 40B. The rapidly rising compliance during the first interval 40A indicates that at least some of the lung opens up at the start of inspiration. This imposes a needless pressure load on the lungs, so the PEEP value set is automatically switched to a higher value. The increase can be performed in specific steps until the all of the first compliance curve 38 is above the first threshold value throughout inspiration. Alternately, a new PEEP can be calculated from measured pressure immediately after compliance exceeds the first threshold value 42.

A second compliance curve 44 is initially constant, but compliance drops below the first threshold value 42 at the end of the second interval 40B. This means that inspiration occurred in the third area 36C in FIG. 2. So excessively high pressure could develop in the lungs. The duration of inspiration is therefore shortened somewhat in order to reduce the risk of harmful excess pressure. A simultaneous change in rate may be necessary to ensure that a sufficient minute volume of breathing gas is supplied to the patient. As an additional safety precaution for some patients, it may be necessary to terminate inspiration as soon as compliance drops below the first threshold value 42. Momentary compliance can also be utilized for determining appropriate reference pressures in pressure-controlled modes, such as PC and PRVC.

The third situation is illustrated by the third compliance curve 46 which is on a constantly high level throughout inspiration. It is even higher than a second threshold value 48. This could mean that the patient's condition has improved and that the first threshold value is no longer relevant. The breathing apparatus can then automatically switch to the use of the second threshold value in the manner described above. Thus, ventilation can be continuously adapted to the patient's condition. If her/his condition worsens, however, a physician should decide on the course of future treatment.

The high compliance of the third compliance curve 46 also suggests that only part of the maximum area 36B is being utilized in ventilation. PEEP can therefore be successively reduced, in order to reduce peak inspiratory pressure (PIP). At the same time, minute volume can be increased by prolongation of inspiration time. If the minute volume is the target volume, the breathing rate can be reduced at the same time.

Momentary compliance needs not be calculated continuously. From the compliance curves shown, it is apparent that the beginning and end of the inspiration are of greatest interest. It would therefore be sufficient for the operation of the method according to the invention to study these parts only.

The invention has been described above in conjunction with a ventilator. But the same method can be implemented in e.g. anesthetic equipment and other

breathing apparatuses. It is also important to avoid the build-up of excessive pressure in healthy lungs, e.g. because more breathing gas is supplied in inspiration than is removed in expiration. Analysis of momentary compliance supplies an additional control parameter for the patient's safety.

Claims

1. A method for controlling a breathing apparatus, a ventilator or anesthetic apparatus in particular, by which method the breathing apparatus is controlled so a breathing gas is delivered to and removed from a respiratory system at the same time as the pressure and flow of the breathing gas is measured, at least when the breathing gas is delivered, characterized in that the respiratory system's momentary compliance is calculated from pressure and flow measured at selected points in an inspiratory phase and compared to a pre-defined threshold value, whereupon one or more of the parameters pressure level, PEEP, the ratio between inspiration time and expiration time and breathing rate is changed if calculated momentary compliance is less than the pre-defined threshold value.
2. A method according to claim 1, characterized in that momentary compliance is calculated virtually continuously throughout the entire inspiratory phase.
3. A method according to claim 1 or 2, characterized in that momentary compliance is calculated as the ratio between momentarily measured flow and the derivative of measured pressure.
4. A method according to any of the above claims, characterized in that PEEP is increased for the next inspiratory phase if calculated momentary compliance is less than the pre-defined threshold value during a first pre-defined part of the inspiratory phase.
5. A method according to claim 4, characterized in that PEEP is reduced for the next inspiratory phase if calculated momentary compliance exceeds a second pre-defined threshold value during the first pre-defined part of the inspiratory phase.
6. A method according to any of the above claims, characterized in that the ratio between inspiration time and expiration time and/or breathing rate is changed if calculated momentary compliance falls below the pre-defined threshold value after a second part of the inspiratory phase expires, so momentary compliance for the next expiratory phase exceeds the pre-defined threshold value during the corresponding part of the inspiratory phase, while a pre-defined minute volume of breathing gas

to the respiratory system is simultaneously maintained.

7. A breathing apparatus (2), a ventilator or anesthetic apparatus in particular, comprising a gas unit (10) for generating a breathing gas pulse which is delivered to a respiratory system (4) during an inspiratory phase, a pressure meter (24, 28) to measure the pressure of supplied gas, a flow meter (26, 30) for measuring the flow of supplied gas and a control unit (12) for controlling the gas unit, characterized in that the control unit (12) is devised to carry out the method according to claims 1-5.
8. A breathing apparatus according to claim 7, characterized in that the pressure meter (28) is arranged in or near the respiratory system (4) to measure the pressure in same, and a flow meter (30) is arranged close to the respiratory system (4) to measure the flow of breathing gas to same.
9. A breathing apparatus according to claim 7, characterized in that the pressure meter (24) and flow meter (26) are arranged in the breathing apparatus (2), and the pressure of the breathing gas in and flow of breathing gas to the respiratory system (4) are calculated in the control unit (12) in the breathing apparatus (2).

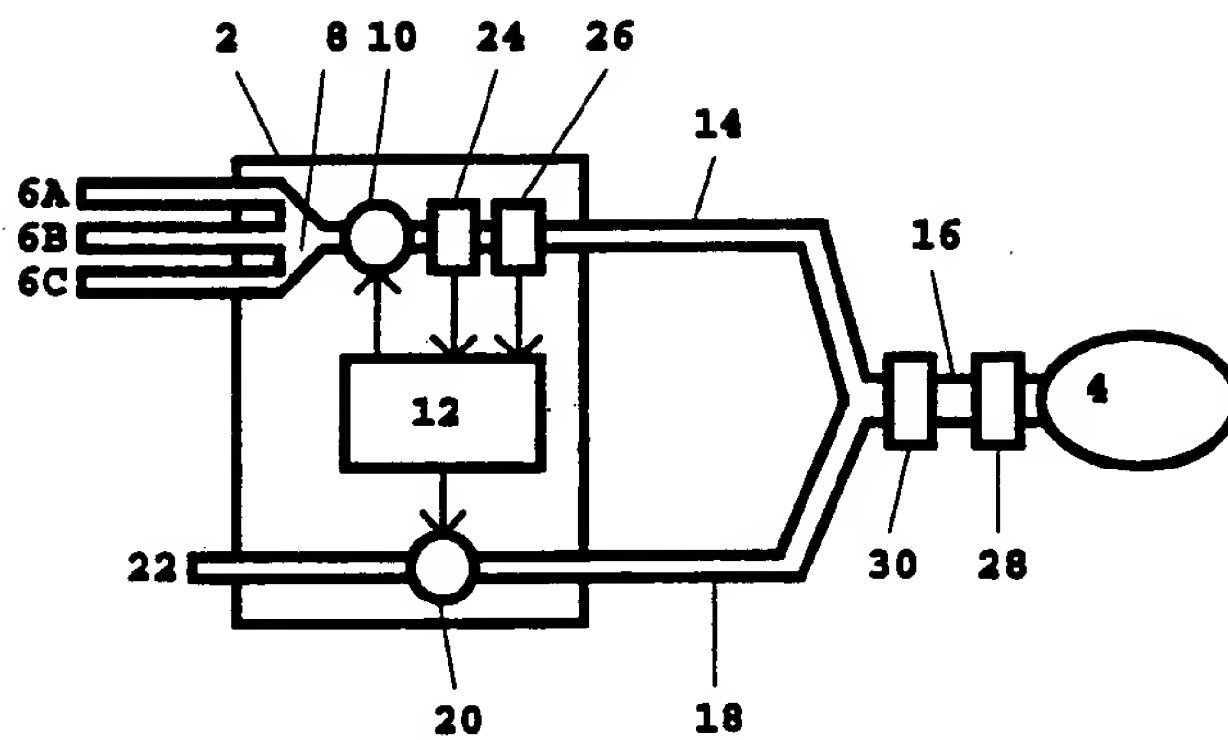


FIG. 1

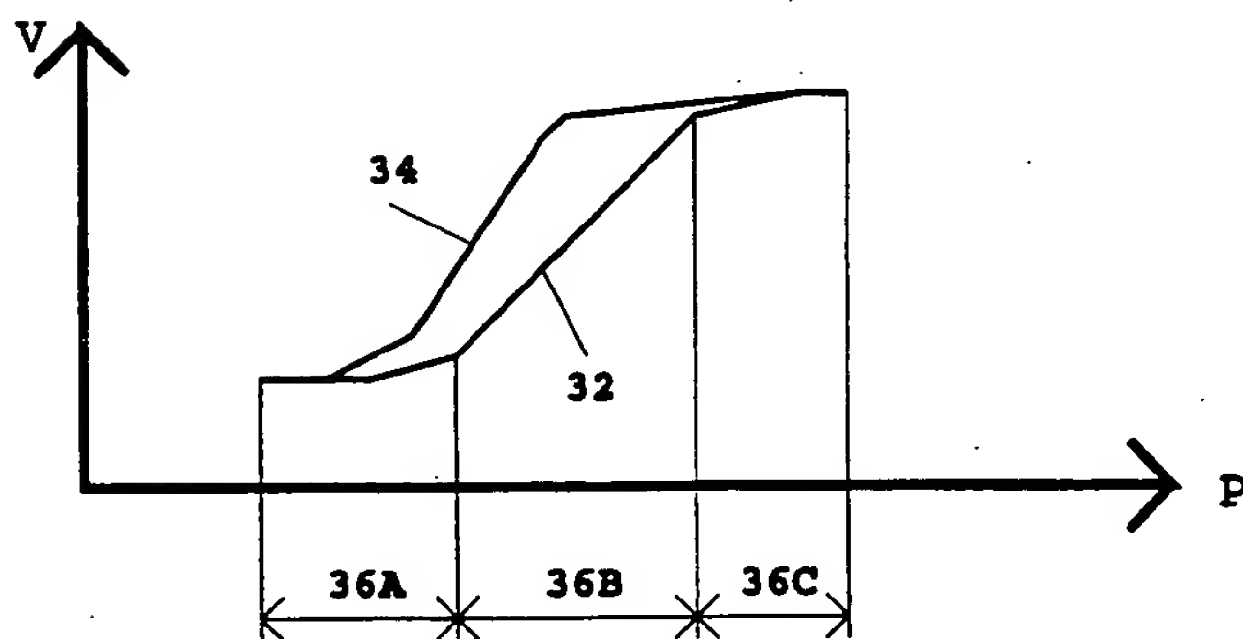


FIG. 2

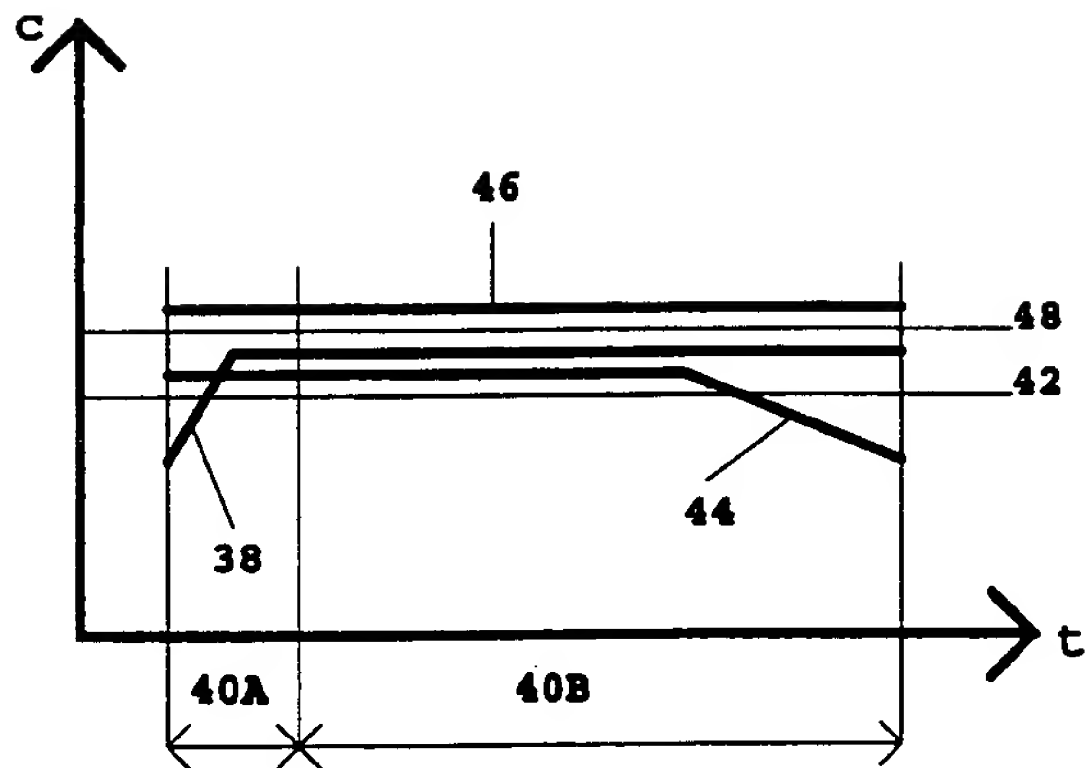


FIG. 3



European Patent
Office

EUROPEAN SEARCH REPORT

Application Number
EP 96 11 6014.0
Page 1

DOCUMENTS CONSIDERED TO BE RELEVANT			
Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int. Cl.6)
A	EP, A1, 0671180 (SIEMENS ELEMA AB), 13 September 1995 (13.09.95) --	1-9	A61M 16/00
A	US, A, 4031885 (J.E.P. DAVIS ET AL), 28 June 1977 (28.06.77) --	1-9	
A	DE, C2, 2744327 (DRÄGENWERK AG), 13 April 1978 (13.04.78) -----	1-9	
			TECHNICAL FIELDS SEARCHED (Int. Cl.6)
			A61M
The present search report has been drawn up for all claims			
Place of search		Date of completion of the search	Examiner
STOCKHOLM		26 February 1997	EVA JOHANSSON
<p>CATEGORY OF CITED DOCUMENTS</p> <p>X : particularly relevant if taken alone Y : particularly relevant if combined with another document of the same category A : technological background O : non-written disclosure P : intermediate document</p> <p>T : theory or principle underlying the invention E : earlier patent document, but published on, or after the filing date D : document cited in the application L : document cited for other reasons & : member of the same patent family, corresponding document</p>			

EPO FORM 1501 03.92 (P0401)